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=> file agricola caplus biossis
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=> s transposase and positive and negative
L1 73 TRANSPOSE AND POSITIVE AND NEGATIVE

```
=> dup rem l1
PROCESSING COMPLETED FOR L1
L2          64 DUP REM L1 (9 DUPLICATES REMOVED)
```

=> s l2 and coda
L3 0 L2 AND CODA

=> d 12 1-10 ti

L2 ANSWER 1 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Genomic insights that advance the species definition for prokaryotes.

L2 ANSWER 2 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI A new insertion sequence, IS14999, from *Corynebacterium glutamicum*.

L2 ANSWER 3 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Simple and straightforward construction of a mouse gene targeting vector
using in vitro transposition reactions.

L2 ANSWER 4 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Method for identification of the indicators of contamination in liquid
samples.

L2 ANSWER 5 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Nucleotide sequence and organization of the multiresistance plasmid pSCFS1
from *Staphylococcus sciuri*.

L2 ANSWER 6 OF 64 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1
TI The positive and negative regulation of Tn10
transposition by IHF is mediated by structurally asymmetric transposon
arms

L2 ANSWER 7 OF 64 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved.

TI (2005) on STN DUPLICATE 2
Germline transformation of the sawfly, *Athalia rosae* (Hymenoptera:
Symphyta), mediated by a piggyBac-derived vector.

L2 ANSWER 8 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Characterization of class 1 integron resistance gene cassettes and the
identification of a novel IS-like element in *Acinetobacter baumannii*.

L2 ANSWER 9 OF 64 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3
TI Long and short mRNAs transcribed from the medaka fish transposon Tol2
respectively exert **positive and negative effects** on
excision

L2 ANSWER 10 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on
STN
TI Identification of genes affecting fluconazole susceptibility in *Candida*
glabrata using a custom transposon.

=> d ab
L3 HAS NO ANSWERS
L1 73 SEA TRANSPOSSASE AND POSITIVE AND NEGATIVE
L2 64 DUP REM L1 (9 DUPLICATES REMOVED)
L3 0 SEA L2 AND CODA

=> d l2 ab
L2 ANSWER 1 OF 64 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
AB To help advance the species definition for prokaryotes, we have compared
the gene content of 70 closely related and fully sequenced bacterial
genomes to identify whether species boundaries exist, and to determine the
role of the organism's ecology on its shared gene content. We found the
average nucleotide identity (ANI) of the shared genes between two strains
to be a robust means to compare genetic relatedness among strains, and
that ANI values of apprxeq94% corresponded to the traditional 70% DNA-DNA
reassociation standard of the current species definition. At the 94% ANI
cutoff, current species includes only moderately homogeneous strains,
e.g., most of the > 4-Mb genomes share only 65-90% of their genes,
apparently as a result of the strains having evolved in different
ecological settings. Furthermore, diagnostic genetic signatures
(boundaries) are evident between groups of strains of the same species,
and the intergroup genetic similarity can be as high as 98-99% ANI,
indicating that justifiable species might be found even among organisms
that are nearly identical at the nucleotide level. Notably, a large
fraction, e.g., up to 65%, of the differences in gene content within
species is associated with bacteriophage and transposase
elements, revealing an important role of these elements during bacterial
speciation. Our findings are consistent with a definition for species
that would include a more homogeneous set of strains than provided by the
current definition and one that considers the ecology of the strains in
addition to their evolutionary distance.

=> s l2 and marker
L4 6 L2 AND MARKER

=> dup rem l4
PROCESSING COMPLETED FOR L4
L5 6 DUP REM L4 (0 DUPLICATES REMOVED)

=> d 1-6 ti
L5 ANSWER 1 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Simple and straightforward construction of a mouse gene targeting vector
using *in vitro* transposition reactions.

L5 ANSWER 2 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Identification of genes affecting fluconazole susceptibility in *Candida*

glabrata using a custom transposon.

L5 ANSWER 3 OF 6 CAPLUS COPYRIGHT 2005 ACS on STN
TI Transposable luciferase expression cassettes for Gram positive
bacteria and their use to monitor bacterial infections by in situ
bioluminescence

L5 ANSWER 4 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI pTn5cat: A Tn5-derived genetic element to facilitate insertion
mutagenesis, promoter probing, physical mapping, cloning, and
marker exchanges in phytopathogenic and other gram-negative
bacteria.

L5 ANSWER 5 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Testing transposable elements as genetic drive mechanisms using Drosophila
P element constructs as a model system.

L5 ANSWER 6 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Identification of IS1356, a new insertion sequence, and its association
with IS402 in epidemic strains of Burkholderia cepacia infecting cystic
fibrosis patients.

=> d ab

L5 ANSWER 1 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
AB In a gene targeting experiment, the generation of a targeting construct
often requires complex DNA manipulations. We developed a set of cassettes
and plasmids useful for creating targeting vectors to modify the mammalian
genome. A positive selection marker
cassette(PGK/EM7p-npt), which included dual prokaryotic and eukaryotic
promoters to permit consecutive selection for recombination in Escherichia
coli and then in mouse embryonic stem cells, was flanked by two FRT-loxP
sequences. The PGK/EM7p-npt cassette was placed between the minimum
regions of a Tn7 transposable element for insertion into another DNA by
means of Tn7 transposase in vitro. We also constructed a
plasmid having a loxP-Zeo-loxP cassette between the modified Tn5 outer
elements. These cassettes can be integrated randomly into a given genomic
DNA through the in vitro transposition reaction, thus producing a
collection of genomic segments flanked by loxP sites (floxed) at various
positions without the use of restriction enzymes and DNA ligase. We
confirmed that this system remarkably reduced the time and labor for the
construction of complex gene targeting vectors.

=> d so

L5 ANSWER 1 OF 6 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
SO Nucleic Acids Research, (05) Vol. 33, No. 5.
CODEN: NARHAD. ISSN: 0305-1048.

=> s (ac or ds) and transpose?

L6 373 (AC OR DS) AND TRANSPOSE?

=> del 16 y

=> s (ac or ds) and transpos?

L6 1795 (AC OR DS) AND TRANSPOS?

=> s 16 and vector

L7 116 L6 AND VECTOR

=> s 17 and transgenic

L8 57 L7 AND TRANSGENIC

=> dup rem 18

PROCESSING COMPLETED FOR L8

L9 41 DUP REM L8 (16 DUPLICATES REMOVED)

=> d 1-10 ti

L9 ANSWER 1 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Transformation of Ac/Ds into rice and Ds
transposition analysis of hybrids

L9 ANSWER 2 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Fumonisin detoxification enzyme gene isolated from environmental
microorganisms, compositions and methods for making fumonisin-resistant
transgenic plants, and detoxification for grains and foods and
feeds

L9 ANSWER 3 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Selection of optimal primers for TAIL-PCR in identifying Ds
flanking sequences from Ac/Ds insertion rice lines

L9 ANSWER 4 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Application of acids transposon system to generate marker gene
free transgenic plants in rice

L9 ANSWER 5 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Method for constructing a tag system comprising transposase
-coding genes and use for tagging plant genes

L9 ANSWER 6 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI GST-MAT vector for the efficient and practical removal of marker
genes from transgenic plants

L9 ANSWER 7 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Methods for site-associated modification of gene activity and nucleic acid
structure

L9 ANSWER 8 OF 41 AGRICOLA Compiled and distributed by the National
Agricultural Library of the Department of Agriculture of the United States
of America. It contains copyrighted materials. All rights reserved.
(2005) on STN DUPLICATE 1
TI Transposon-mediated single-copy gene delivery leads to increased
transgene expression stability in barley.

L9 ANSWER 9 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Timing of transposition of Ac mobile element in
potato.

L9 ANSWER 10 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on
STN
TI Activation of non-autonomous maize transposable element,
Dissociation (Ds), by Ac-transposase in
carrot.

=> d ab

L9 ANSWER 1 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
AB A binary vector CamDs carrying the maize transposon
Ds with activation tagging and gene trap was constructed. The
maize transposon Ac/Ds was transferred into
rice (*Oryza sativa* subsp. *japonica* cv. *Xiushui 11*) by Agrobacterium-
mediated transformation method. The integration of Ac/
Ds into rice genome was confirmed by PCR. The Ds
-inserted transgenic plants were crossed with the
transgenic plants carrying Ac transposase and
a population of 12 hybrids was obtained. One hundred and eight hybrids
consisting of both Ds and Ac were obtained by
resistance assaying. The result of the Basta resistance test indicated
that the excision frequency of Ds element trans-activated by
Ac transposase was 13%, PCR anal. showed the similar
result. The GUS staining indicated that the gene trap system could
capture the expression of the genes in rice genome.

=> d so

L9 ANSWER 1 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
SO Zhongguo Shuidao Kexue (2005), 19(1), 1-6
CODEN: ZSKHBX; ISSN: 1001-7216

=> d 3 agb

'AGB' IS NOT A VALID FORMAT

In a multifile environment, a format can only be used if it is valid in at least one of the files. Refer to file specific help messages or the STNGUIDE file for information on formats available in individual files.

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L9 ANSWER 3 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
AB Infections with the Human papillomavirus (HPV) are related to the development of cervical cancer. It's very important to develop of an HPV prophylactic vaccine. Transgenic plants is a highly-profitable bioreactor, in this experiment, it was planning to establish HPV16-L1 transgenic plants, producing large amount of HPV16-L1 major capsid protein. The HPV16-L1 coding sequence was amplified by PCR with specific primers and plasmid pEGM-HPV16 used as a template, subcloned into middle vector pUCmT and binary vector pBI121 to obtain plant expression Vector pBI-L1. On the T-DNA regions of the pBI-L1 binary vector contained constitutive Cauliflower mosaic virus (CaMV) 35S promoter, nopaline synthase terminator, and neomycin phosphotransferase npt II gene, which allows the selection of transformed plants against kanamycin. The tobacco (*Nicotiana tabacum* L. Crttivar Xanthi) plants were transformed by co-cultivating leaf disks method via *Agrobacterium tumefaciens* LBA4404 harboring the plant expression vector. The regenerated transgenic tobacco plants were selected by kanamycin, and confirmed by PCR, Southern blot and Western blot. PCR and Southern blot analyses confirmed stable integration of the HPV16-L1 gene into the transformed tobacco plants genome. Western blot verified the expressed protein of interest being reactive with the antibody against HPV16-L1, showed that the protein was about 55 kD, consistent with the of HPV16-L1 protein, implying that the given protein was HPV16-L1. The levels of 1.1 expression were up to 0.076 of total soluble tobacco leaf protein by ELISA assay. Expressed protein of transgenic tobacco plants was analyzed by mouse erythrocyte hemagglutination assay (HA) and hemagglutination inhibition assay (HAI), which had the same bio-activity as the natural HPV-16L1 protein, causing murine erythrocyte agglutination and forming VLP by self-assemble in vitro. These results indicate clearly that transgenic HPV16-L1 tobacco plants were generated, and HPV16-L1 protein was expressed effectively in transgenic tobacco plants. This result is an important step close to developing an edible vaccine, which will contribute to the prevention of HPV 16 infectious. Thermal asym. interlaced-PCR, as a PCR-based technique in identifying DNA fragments flanking known sequences, has obtained wide application in different organisms thereby greatly promotes the efficiency in reverse genetics. Unfortunately, in spite of the fact that TAIL-PCR technique has been expanded vastly and adopted in transposon mutagenesis in rice, a reliable, highly reproducible TAIL-PCR procedure especially for rice genomic DNA is still not available, mainly due to the complexity of rice genome and the lack of optimal primers for TAIL-PCR in rice. Given the current situation, we designed 12 specific primers corresponding to 3' end or complimentary to 5' end of Ds insertion, which constitute 32 sets, each with 3 specific primers for three rounds of TAIL-PCR, for screening the optimal combinations of Ds-specific primers. Based on the massive results front pilot expts., two optimal sets of specific primers (Ds3L1/Ds3L2/Ds3S3 at 3' end; Ds5L1/Ds5S3 at 5' end) were chosen, and used together with six arbitrary degenerate (AD) primers, resp., to comparatively investigate the effects of arbitrary degenerate primers on the specificity of TAIL-PCR. Among the tested six AD pruners, AD4 (5'-NTCAGSTWSWGWT-3') possessing 128 fold degeneracy, was proved to

be the most efficient for TAIL-PCR with rice genomic DNA. Moreover, the results also implied that long specific primers in the primary reaction favored the TAIL-PCR by increasing specificity, and different AD primers led significant differences in PCR amplification, presumably due to great difference in degeneracy. The data may provide helpful information for TAIL-PCR technique to improve the efficiency in identifying DNA fragments flanking Ds insertion in rice or other organisms.

=> d 3 so

L9 ANSWER 3 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
SO Shengwu Gongcheng Xuebao (2004), 20(6), 821-826
CODEN: SGXUED; ISSN: 1000-3061

=> d 11-20 ti

L9 ANSWER 11 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Transposon tagging and gene delivery in small grain cereals

L9 ANSWER 12 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Frequency and pattern of transposition of the maize transposable element Ds in transgenic rice plants.

L9 ANSWER 13 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI Improvement of a new transformation method: MAT vector system

L9 ANSWER 14 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Transformation of korean chrysanthemum (Dendranthema zawadskii X D. X grandiflorum) and insertion of the maize autonomous element Ac using Agrobacterium tumefaciens.

L9 ANSWER 15 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 2
TI Germinial virus vector WDV (wheat dwarf virus)-mediated multiple insertions of a maize transposon, Ds (dissociation), in rice

L9 ANSWER 16 OF 41 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved. (2005) on STN DUPLICATE 3
TI Effective selection system for generating marker-free transgenic plants independent of sexual crossing.

L9 ANSWER 17 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Insertion of the maize transposable element Ac into soybean (Glycine max L. Merr.) by Agrobacterium mediated transformation method.

L9 ANSWER 18 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN
TI P gene promoter constructs for floral-tissue preferred gene expression

L9 ANSWER 19 OF 41 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI The transposition frequency of Tag1 elements is increased in transgenic Arabidopsis lines.

L9 ANSWER 20 OF 41 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 4
TI Transposition behavior of the maize transposable element Ac in transgenic haploid tobacco

=> s ((lam e?) or (lam, e?))/au

L10 585 ((LAM E?) OR (LAM, E?))/AU

=> s l10 and transposase
L11 1 L10 AND TRANSPOSASE

=> d ti

L11 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
TI Compositions and methods for targeted gene insertion

=> s l0 and transpos?
L12 1 L0 AND TRANSPOS?

=> d ti

L12 ANSWER 1 OF 1 CAPLUS COPYRIGHT 2005 ACS on STN
TI Preparation and prototropic transposition of some
 γ -acetylenic acids

=> del l12 y

=> s l10 and transpos?
L12 2 L10 AND TRANSPOS?

=> d 1-2 ti

L12 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN
TI Compositions and methods for targeted gene insertion

L12 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2005 ACS on STN
TI From footprint to function: an approach to study gene expression and
regulatory factors in transgenic plants

=> s l10 and (ac or ds)
L13 8 L10 AND (AC OR DS)

=> dup rem l13
PROCESSING COMPLETED FOR L13
L14 5 DUP REM L13 (3 DUPLICATES REMOVED)

=> d 1-5 ti

L14 ANSWER 1 OF 5 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Condylar shape analysis using panoramic radiography units and conventional
tomography.

L14 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1
TI H2O2 induces a transient multi-phase cell cycle arrest in mouse
fibroblasts through modulating cyclin D and p21Cip1 expression

L14 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 2
TI BCR-ABL and interleukin 3 promote hematopoietic cell proliferation and
survival through modulation of cyclin D2 and p27Kip1 expression

L14 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN
TI Compositions and methods for targeted gene insertion

L14 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 3
TI Caspases and programmed cell death in the hypersensitive response of
plants to pathogens

=> d 2 ab

L14 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2005 ACS on STN DUPLICATE 1
AB To defend against the potential damages induced by reactive oxygen
species, proliferating cells enter a transient cell cycle arrest. We

treated mouse fibroblasts with H2O2 and found that sublethal doses of H2O2 induced a transient multi-phase cell cycle arrest at the G1, S, and G2 phases but not the M phase. Western blot anal. demonstrated that this transient cell cycle arrest is associated with the down-regulation of cyclins D1 and D3 and up-regulation of the CKI (cyclin-dependent kinase inhibitor) p21Cip1 expression. We also demonstrate that the induction in p21Cip1 expression by H2O2 is at least partially mediated at the transcriptional level and can occur in the absence of p53 function. Further immunopptn. kinase and immunodepletion assays indicated that in response to H2O2 treatment, the down-regulation of cyclin Ds expression are associated with repression of cyclin D-CDK4, whereas the accumulation of p21Cip1 is responsible for the inhibition of cyclin E and A-CDK2 activity and associated with the down-regulation of cyclin B-CDC2 activity. These data could account for the cell cycle arrest at the G1, S, and G2 phases following H2O2 stimulation. Deletion of p21Cip1, restoration of cyclin D expression, or overexpression of cyclin E alone is insufficient to effectively overcome the cell cycle arrest caused by sublethal doses of H2O2. By contrast, overexpression of the human herpesvirus 8 K cyclin, which can mimic the function of cyclin D and E, is enough to override this transient cell cycle arrest. On the basis of our findings, we propose a model in which moderate levels of H2O2 induce a transient multi-phase cell cycle arrest at least partially through up-regulation of p21Cip1 and down-regulation of cyclin D expression.

=> s l10 and homologous recombination
L15 6 L10 AND HOMOLOGOUS RECOMBINATION

=> dup rem l15
PROCESSING COMPLETED FOR L15
L16 4 DUP REM L15 (2 DUPLICATES REMOVED)

=> d 1-4 ti

L16 ANSWER 1 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
TI Compositions and methods for targeted gene insertion

L16 ANSWER 2 OF 4 BIOSIS COPYRIGHT (c) 2005 The Thomson Corporation on STN
TI Targeted gene insertion in higher plants via homologous recombination.

L16 ANSWER 3 OF 4 CAPLUS COPYRIGHT 2005 ACS on STN
TI Targeted disruption in *Arabidopsis*

L16 ANSWER 4 OF 4 AGRICOLA Compiled and distributed by the National Agricultural Library of the Department of Agriculture of the United States of America. It contains copyrighted materials. All rights reserved.
(2005) on STN DUPLICATE 1
TI Targeted disruption of the TGA3 locus in *Arabidopsis thaliana*.

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DATE: Thursday, August 25, 2005

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DB=PGPB,USPT; PLUR=YES; OP=ADJ

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<input type="checkbox"/>	L7	L6 and activation tagging	1
<input type="checkbox"/>	L6	L3 and homologous recombination	108
<input type="checkbox"/>	L5	L4 and homologous recombination	38
<input type="checkbox"/>	L4	L3 and maize	45
<input type="checkbox"/>	L3	L2 and positive select\$	126
<input type="checkbox"/>	L2	L1 and negative select\$	236
<input type="checkbox"/>	L1	transposase	1183

END OF SEARCH HISTORY